## Amendment to the Claims

- 1. (Amended) A transgenic mouse whose genome comprises a <u>homozygous</u> disruption in anthe endogenous <u>solute carrier family 19 (thiamine transporter)</u>, <u>member 2 (SLC19A2)</u> gene, wherein where the disruption is homozygous said mouse exhibiting, relative to a wild-type control mouse, the transgenic mouse lacks production of functional SLC19A2 protein, and exhibits a reproductive system abnormality.
- 2. (Original) The transgenic mouse of claim 1, wherein the transgenic mouse exhibits a genitourinary system abnormality.
- 3. (Original) The transgenic mouse of claim 2, wherein the transgenic mouse exhibits an abnormality of the testis and epididymus.
- 4. (Original) The transgenic mouse of claim 3, wherein the transgenic mouse exhibits reduced combined testicular and epidiymus weights, relative to a wild-type mouse.
- 5. (Original) The transgenic mouse of claim 3, wherein the transgenic mouse exhibits reduced combined testicular and epididymus weight relative to body weight, compared to a wild-type mouse.
- 6. (Original) The transgenic mouse of claim 3, wherein the transgenic mouse exhibits testicular degeneration.
- 7. (Original) The transgenic mouse of claim 6, wherein the transgenic mouse exhibits degenerative changes of the seminiferous tubules.
- 8. (Original) The transgenic mouse of claim 3, wherein the transgenic mouse exhibits hypospermatogenesis.
- 9. (Original) The transgenic mouse of claim 3, wherein the transgenic mouse exhibits aspermia of the epididymus.
- 10. (Original) A cell or tissue obtained from the transgenic mouse of claim 1.
- 11. (Amended) A transgenic mouse <u>whose genome comprises comprising</u> a heterozygous disruption in an endogenous SLC19A2 gene, wherein the disruption in a homozygous state inhibits production of functional SLC19A2 protein resulting in a transgenic mouse exhibiting a reproductive system abnormality.

Claims 12-13 (Canceled).

- 14. (Withdrawn) A targeting construct comprising:
  - (a) a first polynucleotide sequence homologous to at least a first portion of an endogenous SLC19A2 gene;
  - (b) a second polynucleotide sequence homologous to at least a second portion of the endogenous SLC19A2 gene; and
  - (c) a selectable marker located between the first and second polynucleotide sequences; wherein the targeting construct, when introduced into a murine embryonic stem cell produces a murine embryonic stem cell comprising a disruption in the endogenous SLC19A2 gene.
- 15. (Withdrawn) A murine embryonic stem cell comprising a disruption in an endogenous SLC19A2 gene, the disruption produced using the targeting construct of claim 0.